

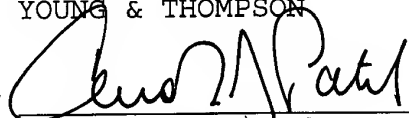
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R E M A R K S

Attached hereto is a marked-up version of the changes made to the claims by the current amendment. The attached page is captioned "VERSION WITH MARKINGS TO SHOW CHANGES MADE."

Respectfully submitted,
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By



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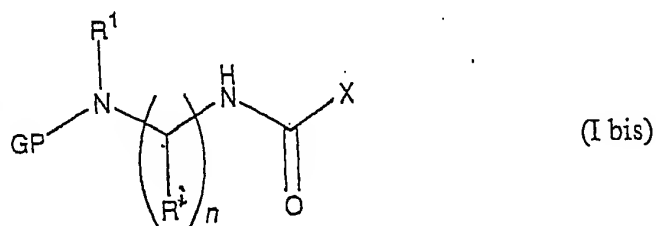
VERSION WITH MARKINGS TO SHOW CHANGES MADE

Amend claim 3 as follows:

--3. (amended) Process according to claim 1-or-2, in which the structure of the activated derivative of carbamic acid is conferred by the N-hydroxysuccinimide group.

Amend claim 9 as follows:

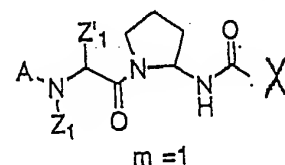
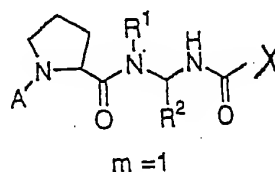
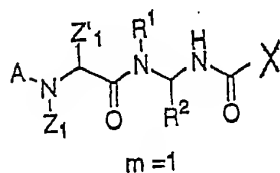
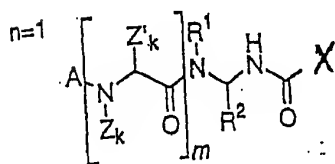
--9. (amended) Compounds according to claim 7-or 8, having the formula (I bis)

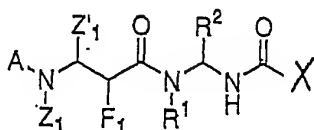
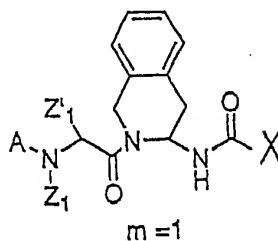
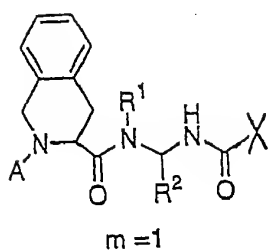
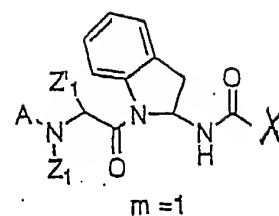
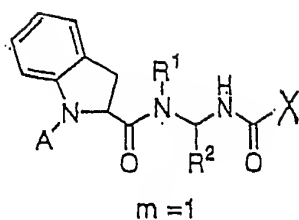


~~in which n, i, GP, X, R¹ and Rⁱ have the meanings given in claims 7 or 8.~~

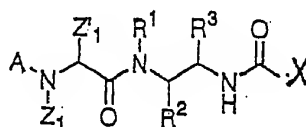
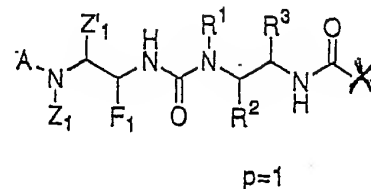
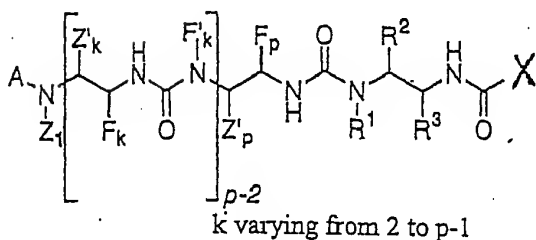
Amend claim 10 as follows:

--10. (amended) Compounds according to claim 7-or 8, having the following formulas:





$n=2$



in which

- m and p are comprised from 1 to 10,
- A represents either a protective group selected from the following groups: a hydrogen atom, an oxycarbonyl (ROCO) group, acyl, alkyl, aryl, urea, phthalimide (with $R1 = \text{ö}$), biotin, or the group A can form with the nitrogen atom with which it is contiguous an NH_2^+ entity,
- Z_k , Z'_k , F_k and F'_k represent independently of each other a hydrogen atom, a protected or unprotected side

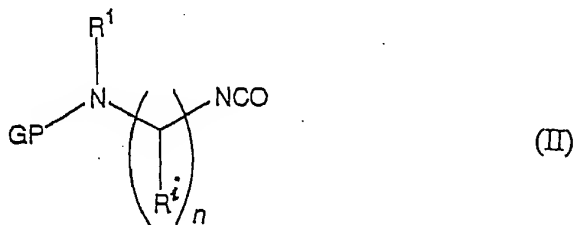
chain of an amino acid selected from natural and synthetic amino acids, a halogen, a (C1-C20) alkyl group, substituted or not, or an aryl group.

Amend claim 12 as follows:

--12. (amended) Compounds according to claim 9, having the formula (Ibis) in which $1 < n < 4$, ~~X is as defined in claim 7~~ and is particularly from p-nitrophenol, N-hydroxysuccinimide, pentafluorophenol, hydroxy-1,2,3-benzotriazole or imidazole, GP is an oxycarbonyl group or acyl group ~~as defined in claim 7~~.

Amend claim 16 as follows:

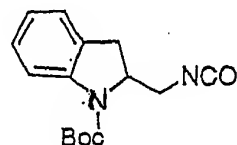
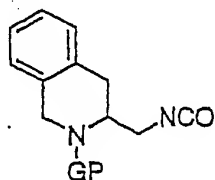
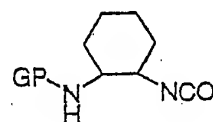
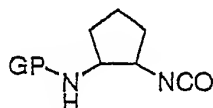
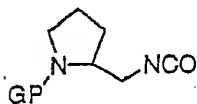
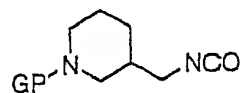
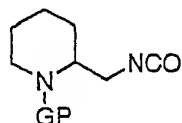
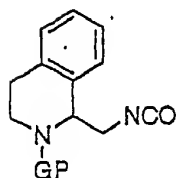
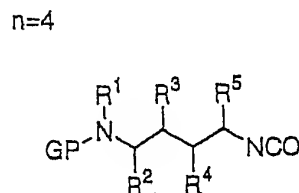
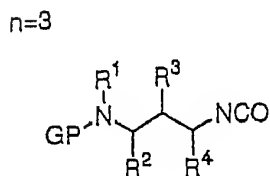
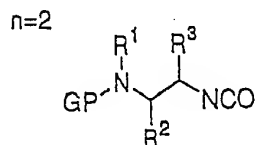
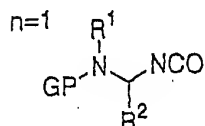
--16. (amended) Compounds according to claim 15, having the formula (II):



~~in which n, i, GP, R1 and Ri have meanings mentioned in claim 15.~~

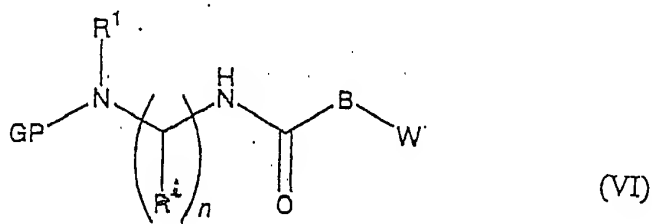
--18. (amended) Compounds according to claim 16, having the formula (II) in which $1 < n < 4$, GP is an oxycar-

bonyl group or acyl group ~~such as defined in claim 15~~, and particularly the following compounds, in particular those for which GP - Boc and Fmoc:



Amend claim 20 as follows:

--20. (amended) Compounds according to claim 19, having the formula (VI):



~~in which n, i, GP, R1, Ri, B and W have the meanings mentioned in claim 19.~~

Amend claim 27 as follows:

--27. (amended) Compounds ~~having one of the formulas (I bis), (II), (III bis), (IV), (VI), (VII)~~ according to ~~one of claims~~ claim 7 to 22, in which the aryl group can be substituted with 1 to 6 substituents selected from: alkyl, alkoxy, amine, ester, urea, amide, carboxylic acid, 1 to 10 carbon atoms, hydroxyl, nitrile, nitro, guanidine, aryl whose cyclic structure contains 5 to 20 carbon atoms, and a halogen atom.

Amend claim 28 as follows:

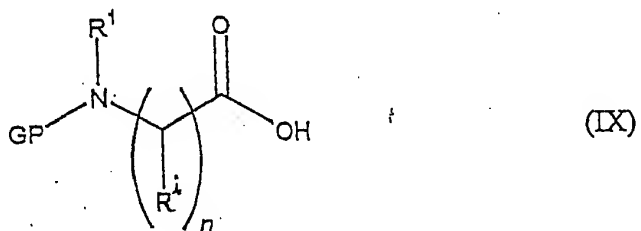
--28. (amended) Compounds according to ~~one of claims~~ claim 7, 15, 19 or 25, in which the alkyl group is substituted with one or several substituents selected from the groups: -COORh, -CONHRh, -COOH, -OH, -ORh, -NHRh, -NH2, -NH(CO)Rh, aryl whose cyclic structure contains 5 to 20 carbon atoms, halogen, carbonyl of 1 to 10 carbon atoms, nitrile, guanidine,

Rh representing an allyl, benzyl, t-butyl, fluorenylmethyl, alkyl group having 1 to 20 carbon atoms, or an aryl group whose cyclic structure contains 5 to 20 carbon atoms.

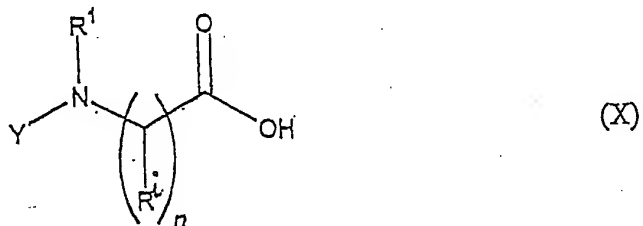
Amend claim 29 as follows:

--29. (amended) Process for preparation according to one of claims 1 to 3, of derivatives corresponding to the formulas (I bis), (II), (III bis) or (IV) according to one of claims claim 7 to 18, from respectively:

- compounds of formula (IX) (for compounds of formula (I bis) and (II))

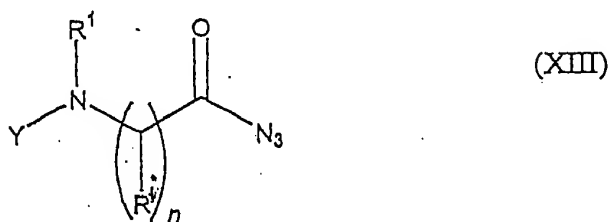
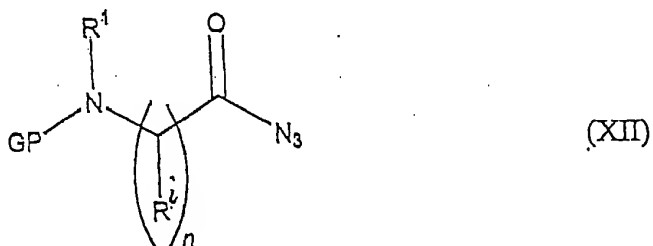


- compounds of formula (X) (for compounds of formula (III bis) and (IV))



comprising

(a) a step of transformation of acid (IX) or (X) into the corresponding acyl azide (XII) or (XIII) respectively



by a suitable treatment,

(b) a step of transformation of acyl azide (XII) or (XIII) by Curtius rearrangement into the corresponding isocyanate (II) or (IV) respectively,

(c) a step of treatment of isocyanate (II) or (IV), preferably not isolated, under conditions permitting obtaining a carbamic acid derivative of formula (I bis) or (III bis).

Amend claim 31 as follows:

--31. (amended) Process for preparation according to claim 1, of compounds of formula (VI) or (VII) according to one of claims claim 19 to 22, comprising the reaction of compounds containing primary or secondary amines or alcohols with one of the products of formula (I bis), (II), (III bis)

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or (IV) ~~according to one of claims 7 to 18,~~ in a solvent, with
or without the addition of an organic or mineral base.

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